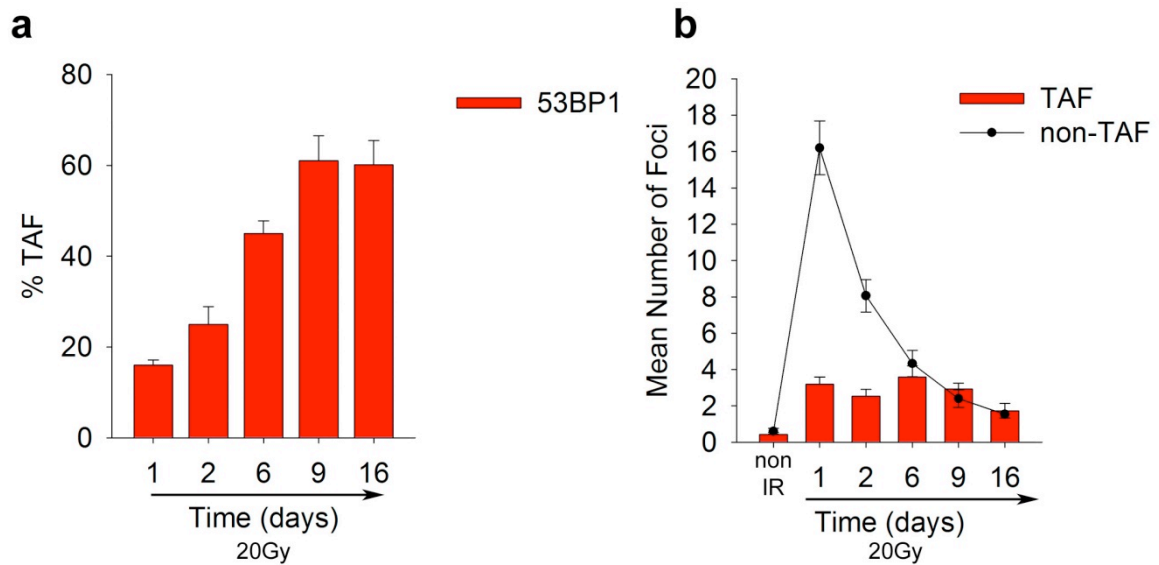
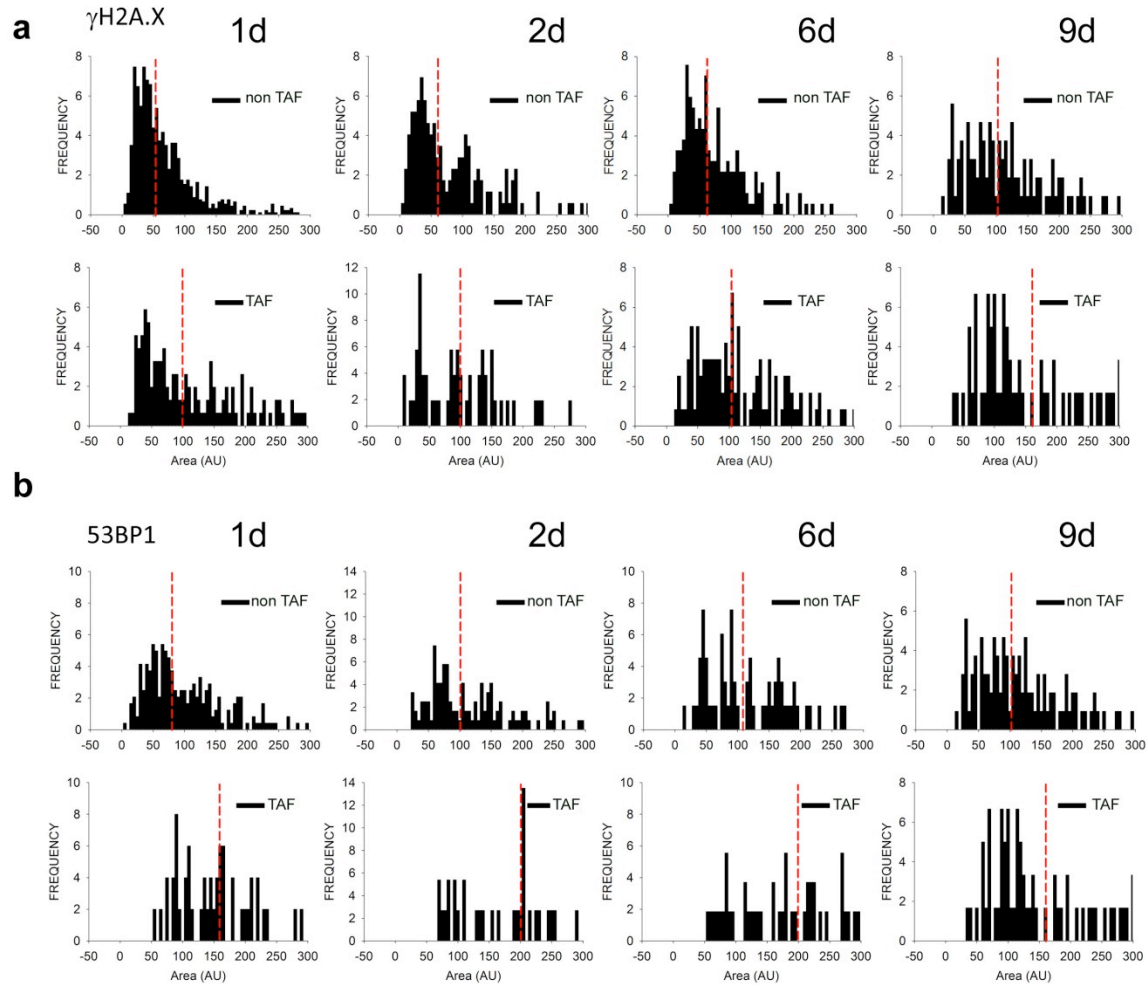


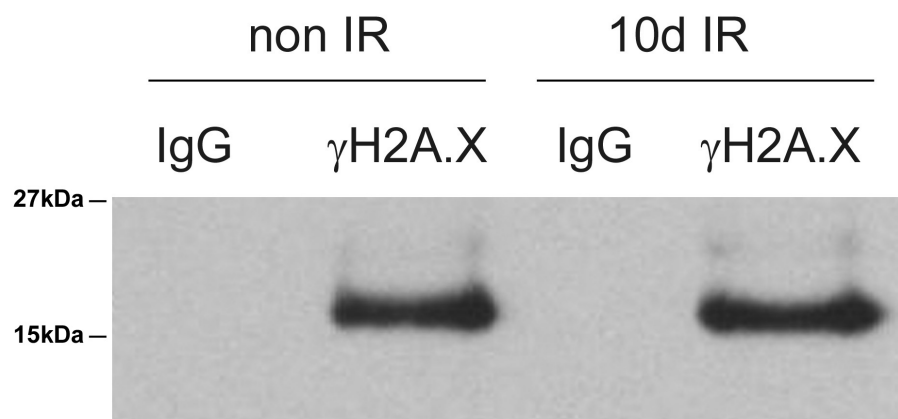
Supplementary information



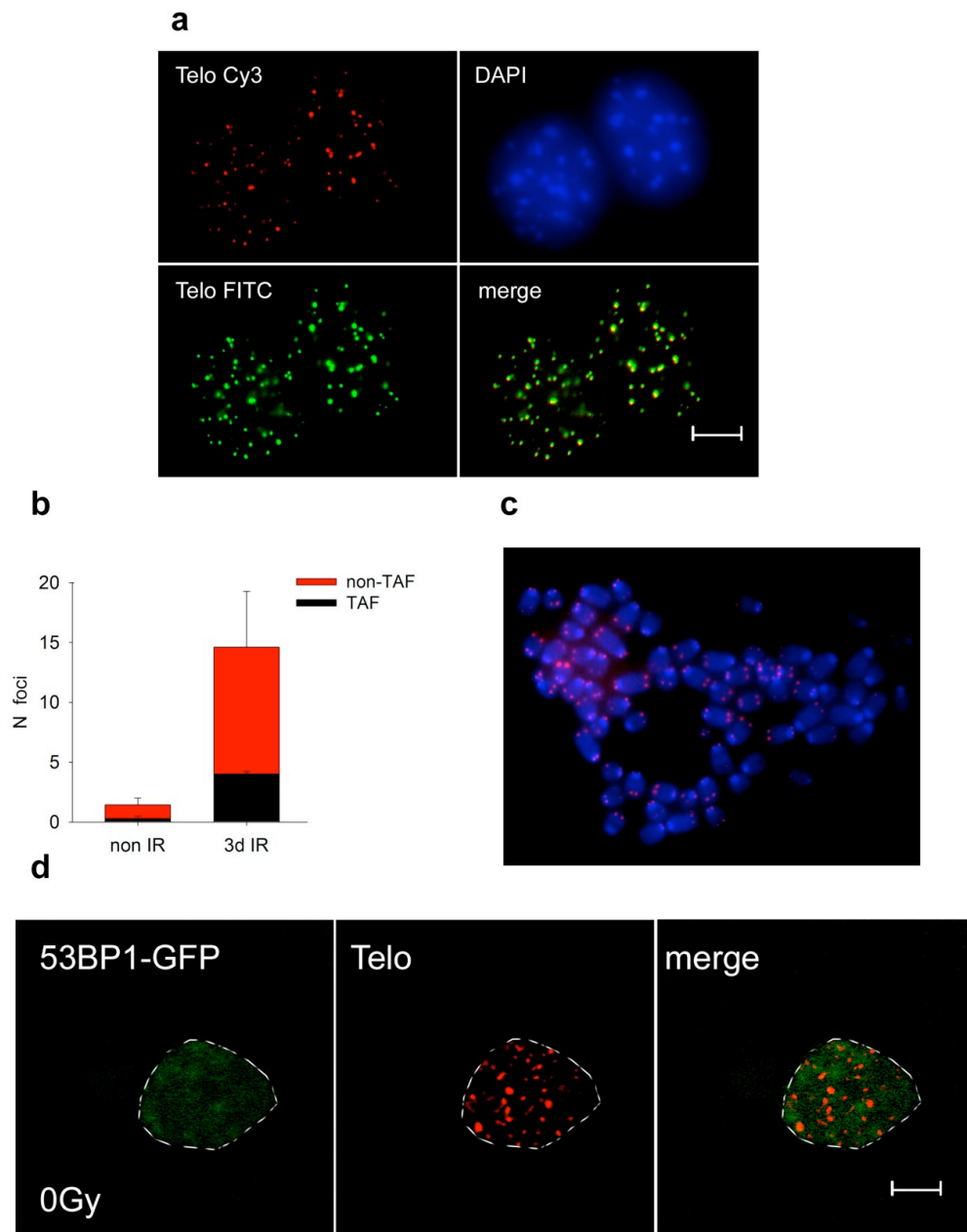
Supplementary Figure S1: Telomere-associated foci (TAF) are persistent following X-ray induced senescence in MRC5 fibroblasts. **(a)** Percentage of 53BP1 foci co-localising with telomeres (%TAF) in MRC5 fibroblasts up to 16 days after Irradiation with 20Gy; **(b)** Mean number of both TAF and non-TAF in MRC5 fibroblasts up to 16 days after Irradiation with 20Gy. Mean \pm S.E.M; n=15.



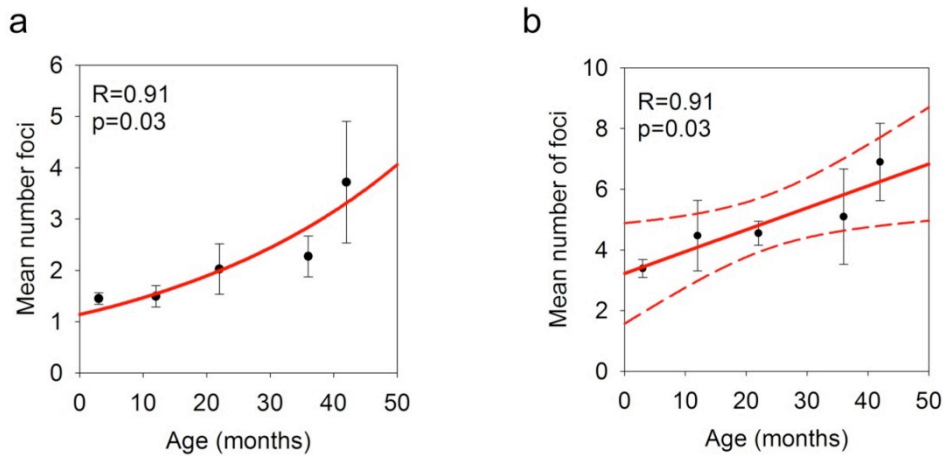
Supplementary Figure S2: Telomere-associated foci (TAF) have larger size than non Telomere-associated foci (non-TAF). Histograms representing distribution of foci size of γ H2A.X (**a**) and 53BP1 (**b**) in MRC5 fibroblasts 1, 2, 6 and 9 days after Irradiation with 20Gy. Dotted red line represents median foci size. Between 100-200 foci were analysed per condition.



Supplementary Figure S3: Extraction of immunoprecipitated γ H2A.X from phenol fractions of both non IR and 10d IR reveals specificity of ChIP assay. Western blot against γ H2A.X in phenol fraction of immunoprecipitated γ H2A.X and IgG in both non IR and 10 days after IR.



Supplementary Figure S4: Microbead mediated telomere PNA probe incorporation does not induce TAF and is telomere specific. (a) After incorporation Cy3 labelled telomere PNA probe in live MEFs, cells were fixed with 2% PFA and hybridised with a FITC labelled telomere probe. Images reveal that majority of red and green spots co-localize Scale bar = 5 μ m; **(b)** Quantification of TAF and non-TAF in non-IR and 3 days following IR in MEFs transfected with AcGFP-53BP1c fusion protein. Data are mean \pm S.E.M n=8; **(c)** After incorporation Cy3 labelled telomere PNA probe in live MEFs, cells were incubated for 2 hours in 10 μ g/ml Colcemid to generate metaphase spreads and then fixed. Image shows that telomere signals can only be found at chromosome ends; **(d)** Mouse embryonic fibroblasts (MEFs) transfected with AcGFP-53BP1c fusion protein show diffuse 53BP1 staining 2 to 8h following microbead incorporation of Cy3 labelled telomere PNA probe; Scale bar = 5 μ m.



Supplementary Figure S5: Mean number of γ H2A.X foci per cell increases with age in both hepatocytes and small intestine crypt enterocytes. **(a)** Mean number of γ H2A.X foci per cell in hepatocytes as a function of age (exponential curve -red- provides best fit, $R=0.91$; $P=0.03$; mean \pm S.E.M of $n=3$ per group); **(b)** Mean number of γ H2A.X foci per cell in small intestinal crypt enterocytes with age (linear curve provides best fit, $R=0.91$; $P=0.03$; mean \pm S.E.M of $n=3$ per group; linear regression-red line, 95% confidence intervals –dotted red lines).